

PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : G03C 1/73, G02B 5/20		A1	(11) International Publication Number: WO 98/13728 (43) International Publication Date: 2 April 1998 (02.04.98)
(21) International Application Number: PCT/US97/11638		(74) Agent: PEOPLES, Veo, Jr.; Peoples & Hale, Suite 500, 1221 Locust Street, St. Louis, MO 63103-2364 (US).	
(22) International Filing Date: 27 June 1997 (27.06.97)		(81) Designated States: CN, FI, JP, KR, SG, European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).	
(30) Priority Data: 08/721,841 27 September 1996 (27.09.96) US		Published <i>With international search report.</i>	
(71) Applicants: BREWER SCIENCE, INC. [US/US]; P.O. Box GG, Rolla, MO 65401 (US). NISSAN CHEMICAL INDUSTRIES, LTD. [JP/JP]; Kowa-Hitotsubashi Building, 7-1, 3-chome, Kanda-Nishiki-cho, Chiyoda-ku, Tokyo 101 (JP).			
(72) Inventors: SABNIS, Ram, W.; 2205 Vichy Road, Rolla, MO 65401 (US). MAYO, Jonathan, W.; 12402 County Road 8010, Rolla, MO 65401 (US). HAYS, Edith, G.; 110 Bryant Road, Rolla, MO 65401-2908 (US). BREWER, Terry, L.; 17971 County Road 8440, Rolla, MO 65401 (US). STRODER, Michael, D.; 3412 South Willow Wick Trail, Springfield, MO 65809 (US). YANAGIMOTO, Akira; 2-24-4, Kichijoji-Hgashi-cho, Musashino-shi, Tokyo 180 (JP). SONE, Yasuhisa; 1-5-17-103, Narashino, Funabashi-shi, Chiba 274 (JP). WATANABE, Yoshitane; 7-17-15-607, Hiraii, Edogawaku, Tokyo 132 (JP). EMA, Kiyomi; 5-38-24, Nishitsuga, Wakaba-ku, Chiba-shi, Chiba 264 (JP).			
(54) Title: ULTRA THIN ORGANIC BLACK MATRIX			
(57) Abstract			
<p>An organic black matrix having high resistivity ($\geq 10^{11}$ ohm/square), high optical density (≥ 2.0) at ultra thin film thicknesses (≤ 1.0 microns) for improved STN and TFT pixel display applications is made possible by combining polyimide/dye solutions and mixed metal oxide pigment dispersions at a weight/weight ratio of dye to pigment of 1:15 to 3:15. The need for low resistivity carbon black, as a replacement for sputtered chrome, is negated.</p>			
<pre> graph TD A[PREPARE SUBSTRATE] --> B[ADHESION PROMOTER] B --> C[COAT BM] C --> D["α-BAKE"] D --> E["β-BAKE"] E --> F[COAT PHOTORESIST] F --> G[EXPOSE] G --> H[DEVELOP] H --> I[STRIP PHOTORESIST] I --> J[FINAL CURE OF BM] J --> K[APPLY OTHER COLORS] </pre>			

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	IS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	MN	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MR	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MW	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MX	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	NE	Mexico	US	United States of America
CA	Canada	IT	Italy	NL	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NO	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	PL	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	PT	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	RO	Poland		
CM	Cameroon	KR	Republic of Korea	RU	Portugal		
CN	China	KZ	Kazakhstan	SD	Romania		
CU	Cuba	LC	Saint Lucia	SK	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SG	Sodan		
DE	Germany	LK	Sri Lanka		Sweden		
DK	Denmark	LR	Liberia		Singapore		
EE	Estonia						

ULTRA THIN ORGANIC BLACK MATRIXBackground of InventionField of Invention.

The present invention relates to organic black matrices for color filter plate manufacture and to methods for making the same. It particularly relates to nonconductive black matrices, having very high resistivity, and having high optical density at ultra thin film thicknesses.

Background of the Prior Art.

10 Multicolor liquid crystal displays (LCDs) are routinely produced having a thin, light-absorbent film, called a black matrix, applied to an array of color pixels which together form a color filter plate. The processing of such color filter plates remains one of the most troublesome steps during 15 mass production of the LCDs because they use *sputtered chrome* black matrices.

A spincoatable, organic polymer based black matrix would tend to be more environmentally friendly than chrome, easier to reproduce, and offer lithographic processing advantages. 20 However, there are at least two types of color filter plates for LCDs where organic polymer black matrices are woefully expensive and/or lack the desired performance. That is, the unavailability of an organic black matrix photoresist having sufficiently high resistivity and sufficiently high optical 25 density at ultra thin film thickness, has hampered the advancement of 1) thin film transistor (TFT)-arrays for TFT-LCDs and 2) super twisted nematic (STN) LCDs.

For purposes of illustration, Figures A, A(a), A(b), and A(c) show a somewhat oversimplified stepwise illustration of the coating and imaging of organic black matrix photoresists onto either TFT or STN color pixels for large area LCDs. The 5 RGB (red, green, blue) color pixels 1 could be either TFTs or STNs. Such color filter plates have an Indium-tin-oxide (ITO) electrode layer or layers 2 and signal lines 3. All are normally applied to a glass substrate 4. The illustration shows the exposure of UV light through the glass substrate for 10 purposes of lithographically imaging the organic black matrix 5.

In these TFT and STN applications, high resistivity (so as to be nonconductive) is mandatory in order to avoid electrical coupling between the ITO electrode layer and the signal 15 lines. Otherwise, coupling with the signal lines (sometimes referred to as data lines) causes vertical crosstalk.

The optical density (O.D.) of the black matrix must be greater than 2.0 in order to block the transmission of light to the TFT or STN displays. Otherwise, photo leaks from non- 20 display areas will reduce the contrast ratio and create adverse photo leakage current. In other words, one can enhance the contrast of the LCD by eliminating the light leakage which would otherwise occur through spaces patterned between the red, green, and blue (RGB) pixels on the color 25 filter plate. The technical goal is to keep the light transmission at or below 1%, across the entire spectrum of from ultraviolet to infrared, at ultra thin black matrix film thicknesses.

It has proven extremely difficult, if not impossible, to manufacture a nonconductive organic black matrix having an O.D. greater than 2.0 at ultra thin black matrix film thicknesses. Although O.D. greater than, or equal to, 2.0 have been achieved for organic black matrices at a 2 micron polymer thickness, such layer thicknesses are at the threshold for a number of defects. For example, the so-called reverse tilt inside each pixel display area occurs at 2 microns. Reverse tilt causes after-image and contrast deterioration. Overcoming that drawback, *inter alia*, requires a nonconductive, organic black matrix film having O.D. greater than 2.0 at a thickness of 1.0 micron or less.

Despite the high cost-of-ownership, a complex manufacturing process (sputtering), and potential environmental problems, and despite a higher reflectance than desired, the most common black matrix material has continued to be sputtered chrome, rather than spin-coated organic polymers. Although vacuum evaporation and other coating techniques for metals such as nickel and aluminum and even chromium have been devised, sputtered chrome remains the most common technique and material because other techniques and other materials lack sufficiently high O.D. (>2.0) to provide enhanced contrast and high resolution, at sufficiently ultra thin thicknesses (1 micron or less) and at sufficiently high resistivity (at least 10^5 ohm/square) to be commercially effective for STN and TFT application.

For example, Latham's U.S. patent 4,822,718 disclosed in 1989 potential black matrices of the organic type made from

polyamic acid/dye combinations (as distinguished from pigment-dispersions). The resistivity of these light-absorbing layers was found to be as high as 3.0×10^{15} ohm/square.

It has been disclosed by Hessler, et al., in the article 5 "Pigment-Dispersed Organic Black-Matrix", SID Digest, 26:446, (1995), however, that Latham's mixtures of the red and blue dyes bound in polyimide composition as well as mixtures of red and blue pigments in pigment-dispersed organic black matrices disclosed by others did not come close to the desired O.D. 10 (greater than 2.0), even when coated at film thicknesses greater than 3 microns. Changing the ratios of red to blue pigments, or adding low levels of violet, yellow or green pigments to the red and blue pigment mixtures did not improve overall O.D. performance of the formulations. Carbon black 15 pigment, dispersed in acrylic polymer, did, however, achieve 0.01% light transmission for some samples. An average O.D. of 2.8 for a 1.5 micron, spin-coated film, over the whole spectrum of 400-700 nm, was made possible by a critical selection of dispersants. However, the resistivity was a dis- 20 appointing 90K ohm/ square.

As disclosed by Yamanaka in his article, "Integrated Black Matrix on TFT Arrays", SID Digest, 23:789 (1992), carbon black, even with an advanced acrylic photo polymer, does not achieve O.D. greater than 2.0, at thicknesses less than 25 2 microns without sacrificing resistivity and creating cross-talk. Yamanaka also describes the disadvantage of a 2 micron "step size" (or thickness). It is so large that it results in reverse tilt.

U.S. patent 5,368,976, by Tajima, et al., discloses another example of pigment-dispersed color filter compositions. An alkali-soluble block copolymer is employed as a binder for a radiation-sensitive compound and pigments such as pigment Black 1 and pigment Black 7 but with unsuitably large particle sizes (filtered at 10 microns). Although it is well known that pigment Black 1 does not provide the requisite O.D. greater than 2.0 at practical thicknesses, pigment Black 7 (better known as simply "Carbon Black") does provide O.D. greater than 2.0 as shown by Hessler, et al. Yet, as Yamanaka explains, even at thicknesses less than 2.0 microns, the threat of crosstalk exists when the black matrix is too conductive for effective STN or TFT applications, and particle sizes of 10 micron diameter would lead to films having reverse tilt.

Suginoya, et al., in their article "Self-Alignment Fabrication of the ITO Electrode Pattern on an Electro-deposited Tricolor Filter in a Black Matrix: An Application to STN-LCDs", Proc. of SID, 32:201, (1991), identify another shortcoming of STN-LCDs having organic black matrices made from carbon black. That is, although the green filter of the display provides a good shutter which transmits less than 1% of incident light at 365, 405 and 435 nm, the red filter does not. It has transmittance of 4% at 365 nm and 6% at 405 nm. Also, the blue filter is even worse, having transmittance of 35% at 405 nm and of 55% at 435 nm. A separate approach was therefore necessary for light above 400 nm than for light below 400 nm. The black matrix average transmittance was sub-

stantially greater than 1.0% throughout the spectrum, when using this complicated approach (Fig. 13 of the reference). It was closer to 10%.

Accordingly, neither the past dye-based nor the past 5 pigment-based organic black matrices could effectively provide O.D. greater than 2.0, at thicknesses less than 1 micron without disaffecting STN and TFT performance. Furthermore, although the dye-based black matrices have the requisite resistivity to provide improved avoidance of crosstalk from 10 electrical coupling between ITO electrodes and signal lines, the substitution of sufficient dye mixtures for a portion of the pigment-dispersed material needed to raise its resistivity beyond 10^5 ohm/square would be expected to weaken the O.D. to below 2.0 unless a film thickness greater than 2.0 microns was 15 applied to the pixels and substrate. Only Carbon Black (Pigment Black 7) with its low resistivity of 90×10^2 ohm/square comes close to O.D. 2.0 at 1.5 micron thickness.

It would therefore be highly unexpected that any combination of dye-based and pigment-based organic black matrices, 20 other than Carbon Black, would provide improved O.D. of over 2.0, without requiring a film thicker than 2.0 microns and Carbon Black is too conductive for effective STN and TFT application. The vast difference in resistivity of carbon black pigment versus various organic dyes would lead one to 25 believe that any appreciable amount of such pigment added to the dyes in order to effectively increase O.D. beyond 2.0 would be significantly detrimental to the resistivity per unit volume of the final material. Yet other pigments have had too

low an O.D. to even be considered for ultra thin black matrices.

It is an object of the present invention to provide a stable organic black matrix having O.D. greater than 2.0, at 5 thicknesses less than 1 micron but surface resistivity greater than 10^5 ohm/square.

SUMMARY OF THE INVENTION

The present invention fulfills the above-stated objective and others by a synergistic composition of polyimide-dye-pigment. The black matrix of this invention has high resistivity and an O.D. greater than 2.0 at a film thickness of 10 1 micron or less.

The material patterns on various display applications have excellent evenness, high strength, superior environmental 15 safety, high stability, good shelf-life and a low cost of production.

BRIEF DESCRIPTION OF THE DRAWINGS

Figures A, A(a), A(b), and A(c) illustrate coating and 20 imaging organic black matrices photoresist onto generally STN or TFT color pixels for large area LCDs.

Figure 1 is a flow chart explaining the manufacturing process of a black matrix according to the present invention.

Figure 2 is a graph showing transmission spectra of 1 μm cured film obtained in Example 1.

25 Figure 3 shows sets of lithography data using both convection oven and hot plate beta-bake processes.

Figure 4 is a schematic diagram illustrating photo of a resolution dagger.

Figure 5 demonstrates Scanning Electron Microscope (SEM) of the black matrix material after resist removal and final 5 cure.

Figure 6 is a schematic diagram illustrating typical surface roughness measurement of the cured film.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

The present invention fulfills the above-stated objective 10 and others by a synergistic composition of polyimide-dye-pigment black matrix mixtures. Such black matrix compositions preferably comprise, by weight, 4% to 8% polyimide binder, 9% to 13% colorant (dye plus pigment), 80% to 85% solvent and 0.3% to .8% dispersant. The weight/weight ratio of dye: 15 pigment will range from about 1:15 to about 3:15, preferably 2:15. Surprisingly, the combined O.D. of organic dyes plus non-carbon black pigments is greater than the sum of their parts at less than 1.0 micron (hereinafter referred to as ultra thin) film thickness.

20

POLYIMIDE/DYE

The polymer vehicles, for the black matrix compositions of the present invention, will typically include polyimide precursors which react *in situ* to form polyimide resins. Preferably, the precursors will consist of a polyamic acid 25 prepared by reacting oxydianiline (ODA) with pyromellitic dianhydride (PMDA) or by reacting ODA with PMDA and/or benzo-phenone tetracarboxylic dianhydride (BTDA). These components

are typically employed in approximate stoichiometric amounts. Other suitable polyamic acids and polyamic precursors typically employed for the polyimide vehicles of this invention are conventional components as, for example, those listed in 5 Table 1 below. Note that it is possible to include water soluble polymers such as polyvinyl/pyrrolidone and other conventional resins such as novolac in admixture with these precursor components without departing from the Applicants' definition of polyimide precursors.

10

TABLE 1

	Diamines	Dianhydrides
15	p-Phenylenediamine tetracarboxylic	3,3',4,4'-Benzophenone dianhydride
20	3,3'-dimethyl-4,4'-diaminobiphenyl	Pyromellitic dianhydride
25	1,4-bis(4-aminophenoxy)benzene tetracarboxylic	3,3',4,4'-Biphenyl dianhydride
30	4,4'-Bis(4-aminophenoxy)-biphenyl	4,4'-Oxydiphthalic anhydride
	Bis-4(-[4-aminophenoxy]phenyl)ether	3,3',4,4'-Diphenylsulfone tetracarboxylic-dianhydride
	4,4'-Oxydianiline	
	4,4'-Diaminodiphenyl sulfide	
	4,4'-Diaminodiphenyl sulfone	
	2,2-Bis(4-[4-aminophenoxy]phenyl)sulfone	
	9,10-bis(4-aminophenyl)-anthracene	

The dye components of the present invention are soluble organic dye combinations which can effectively absorb light over the broad spectrum of infrared to ultraviolet. It is critical that the dyes be soluble in solvents which dissolve the polyimide vehicle (which solvents will be discussed later). In a particularly preferred embodiment of the present

invention, a weight/weight ratio of from about 1:3 to 1:5 of Orasol brown 6RL (Solvent brown 44) to Orasol blue GN (Solvent blue 67) provides a surprisingly effective component for this invention. Such a mixture is commercially available under the 5 trade name DARC-100 sold by Brewer Science, Inc.

Other combinations of Blue, Red, Orange, etc., dyes capable of producing high resistivity black coatings in U.S. patent 4,822,718 may be employed.

In a preferred embodiment of the present invention, all 10 of the dye component together with a substantial amount of the polyimide component (at least 50%) may be provided in accordance with the black coating compositions disclosed in U.S. patent 4,822,718, which patent is herein incorporated-by-reference to this disclosure.

15 Highly preferable dyes for the present invention may also include Solvent Black 35 (Zapon Black X50, BASF), Solvent Black 27 (Zapon Black X51, BASF), Solvent Black 3 (Neptun Black X60, BASF), Solvent Black 5 (Flexo Black X12, BASF), Solvent Black 7 (Neptun Black NB X14, BASF), Solvent Black 46 20 (Neptun A Black X17, BASF), Solvent Black 47 (Neopin Black X58, BASF), Solvent Black 28 (Orasol Black CN, Ciba-Geigy), Solvent Black 29 (Orasol Black RL, Ciba-Geigy), and Solvent Black 45 (Saviny Black RLS, Sandoz Corp.).

PIGMENT-DISPERSIONS

25 One surprising aspect of the present invention is the ability to obtain optical density > 2.0 at ultra thin film thicknesses without having to resort to the use of Carbon Black.

Although the pigments employed in this invention are indeed macroscopic particles having high surface area and a strong tendency to flocculate which can create haze and light scattering aggregates or variations in rheology, when the 5 pigments of this invention are suitably dispersed and admixed with the polyimide/dye solutions, excellent evenness is achieved.

The pigments of this invention are mixtures of inorganic metal oxides or mixed metal oxides selected from metal oxides 10 of copper, manganese, chromium, iron, magnesium, aluminum, tin, zinc, titan, nickel, cobalt and mixtures thereof. These metal oxides are preferred for purposes of forming the so-called spinel structure. They unexpectedly provide a synergy 15 that together with dyes of this invention will achieve O.D. > 2.0 at ultra thin film thicknesses, while their combined surface resistivity is magnitudes above the 10^5 ohm/square threshold required for effective STN and TFT application.

Suitable pigments may include pigment Black 22 (color index (C.I.) 77429), such as Fast Black 100 sold by Bayer 20 Corp.; Pigment Black 26 (C.I. 77494) such as Daipyroxide TM Black 3550 and 3551 sold by Dainichiseika Color & Chemical Manufacturing Co., Ltd. of Japan; Pigment Black 27 (C.I. 77502); Pigment Black 28 (C.I. 77428) sold as Harshaw 9875 M Plus by Engelhard Corporation; and also mixtures of Pigment 25 Green 50 (C.I. 77377) with Pigment Blue (C.I. 77346), and Pigment Red (C.I. 77491). It is particularly preferred to use Pigment Black 26 (C.I. 77494) composed of mixed metal oxide of copper, manganese and iron.

It is preferred in manufacturing the pigment dispersions of this invention to employ as the primary particle size of the pigments less than 100 nanometers (nm), particularly less than 50 nm. The particles of these pigments may also be 5 coated with inorganic layers of silica, alumina, or zirconia for purposes of improving the stability of the dispersions.

Dispersants, effective for a Newtonian dispersion of the pigment in the polyimide vehicle and solvent system, were desired. Cationic dispersants are preferred, particularly 10 cationic dispersants composed of solutions with high molecular weight blocked copolymers with chemical groups having strong affinity to the pigments of this invention. One such particularly preferred dispersant is that sold under the trademark Disperbyk-163 by Byk-Chemie. Other such cationics 15 include, for example, the trade names Disperbyk-160, 161, 162, 164 and 166. Anionic and nonionic dispersants are also suitable. A list of such dispersants may be seen in Table 2 below.

TABLE 2

	<u>Dispersing Agent</u>	<u>Company</u>	<u>Ionic Nature</u>
5	Disperbyk-160	Byk-Chemie	Cationic
	Disperbyk-161	Byk-Chemie	Cationic
	Disperbyk-162	Byk-Chemie	Cationic
	Disperbyk-163	Byk-Chemie	Cationic
	Disperbyk-164	Byk-Chemie	Cationic
	Disperbyk-166	Byk-Chemie	Cationic
10	Lactimon	Byk-Chemie	Anionic
	Bykumen	Byk-Chemie	Anionic
	Dumasperse 535	Hickson	Anionic
	Dumasperse 540	Hickson	Anionic
	Dumasperse 545	Hickson	Anionic
	Mazsperse 85B	PPG	Nonionic
15	Mazsperse SF 19	PPG	Nonionic
	Nuosperse 657	Huls	Nonionic
	Nuosperse 700	Hul	Anionic
	Solsperse 12000	Zeneca	Anionic
20	Solsperse 27000	Zeneca	Nonionic

The dispersion is carried out in, for example, Eiger Mini-100 motormills using suitable beads, e.g., 0.65 mm yttrium stabilized zirconium oxide beads.

Preferably, the pigments are dispersed in a Newtonian dispersion having dispersant in an amount comprising 5% the weight of the pigment, a solution of polyimide precursor (22.7% solids in solution) and a substantial amount of solvent.

SOLVENTS

30 A solvent effective for solubilizing the polyimide vehicle and dye compound is selected. The most preferred solvents for the polyimide/dye component and for the pigment dispersion are N-methyl-2- pyrrolidone (NMP) and cyclohexanone. Other suitable solvents may include dimethylacetamide (DMAc), N,N-dimethylformamide (DMF), bis-2-methylethyl ether (diglyme), tetrahydrofurfuryl alcohol (THFA),

dimethylsulfoxide (DMSO), xylenes, cyclic ketones, alcohols, esters, ethers and mixtures thereof. A polymer/dye solution of such dyes shall also be suitable for the effective pigment dispersion, previously discussed.

5 The black matrix compositions of this invention (1) can be applied to substrates by spin coating, (2) can be imaged using the same process employed to image the RGB pixels, (3) developed using the same development process as the pixels, and (4) provide an average O.D. for 1 micron film 10 thicknesses over the region 400-700 nm of 2.4 O.D. The shelf life is excellent, i.e., three months in the freezer or three weeks at room temperature. The surface resistivity is very high, i.e., in the range of 10^{11} ohm/square, and cost of production is much lower than sputtered chrome black matrices.

15 EXAMPLE 1

In a plastic beaker is added 11.6 g of N-methyl-pyrrolidone (NMP), 82 g of cyclohexanone, 1.5 g of disperbyk-163, 20 g of DARC polymer and 30 g of Pigment Black 26. The mixture is stirred with a spatula for about 20 5 minutes till homogeneous. This pigment slurry is then added to Eiger M-100 mill at 1000 rpm over a period of 15 minutes, using 0.65 mm yttrium stabilized zirconium oxide beads. The beaker is then rinsed with 20 g of cyclohexanone and rinse added to the mill. The grind speed is slowly increased to 25 3000 rpm. The pigment is ground at this speed for 2 hours. In a separate plastic beaker equipped with mechanical stirrer is added 32.4 g of DARC polymer (22.9% in NMP only) and 60 g of DARC 100. The mixture is stirred for 10 minutes. After

2 hours, the mixture of DARC polymer and DARC 100 is added to the mill at 1000 rpm. DARC polymer is synthesized by dissolving 48.1 g of oxydianiline in NMP, then added 51.8 g of pyromellitic dianhydride and reacted for 5 hours at 40°C. The 5 beaker is rinsed with 50 g of N-methyl-pyrrolidone and the rinse is added to the mill at 1000 rpm. The mill is then run at 3000 rpm for 90 minutes. Formulation is then filtered through 0.2 μ m filters. The formulation is summarized in Table 3. A resistivity of 5.6×10^{11} ohm/square, at a thickness of 1 micron and an O.D. of 2.4 were measured.

TABLE 3

Chemical Constituent	Weight in grams
15 N-methyl-pyrrolidone	61.6 g
Cyclohexanone	102 g
Disperbyk-163	1.5 g (5% of the weight of pigment)
DARC polymer	52.4 g (22.9% in NMP only)
Pigment Black 26	30 g
20 DARC 100	60 g

Table 4 illustrates preferred composition with reference to Weight % for 2.4 optical density and 10^{11} ohm/square surface resistivity, at 1 micron film thickness.

TABLE 4

Chemical Composition	Weight %	Best Mode
5 Polyamic acid	4% ~ 8%	5.7%
Colorant (dye + pigment)	9% ~ 13%	11.5%
Dispersant	0.3% ~ 0.8%	0.8%
Solvent	80% ~ 85%	82.0%

10

EXAMPLE 2

In a plastic beaker is added 375 g of NMP, 375 g of cyclohexanone and 250 g of Pigment Black 26 (Daipyroxide TM Black 3551) with the primary particle size ranging from 10 to 20 nm and the surface of which is covered with thin silica layers. The mixture is stirred with a spatula for 5 minutes till homogeneous. This pigment slurry is then added to a ball mill together with quartz beads with 5 mm diameter of particle size, and ground at the speed of 100 ~ 200 rpm for 2 weeks. This mixture is dispersed at the speed of 7,000 rpm for 20 5 minutes in a homogenizer (Nihon Seiki Kaisha Ltd.) by adding 47.8 g of NMP and 21.0 g of cyclohexanone to it. In a separate glass beaker equipped with a mechanical stirrer is added 67.4 g of DARC polymer (20 wt % in NMP only), 54.8 g of DARC 100, 69.8 g of NMP and 33.5 g of cyclohexanone. The 25 mixture is dispersed in a homogenizer at the speed of 7,000 rpm for 5 minutes. In a homogenizer is added 225.3 g of the polymer/dye mixture and 175.7 g of the black pigment slurry. The homogenizer is then run at 10,000 rpm for 10 minutes. Formulation is then filtered through 0.2 μ m 30 filters. The formulation is summarized in Table 5.

TABLE 5

Chemical Composition	Weight %
5 Polyamic acid	4% ~ 8%
Colorant	6% ~ 10%
Dispersant	0.1% ~ 0.4%
Solvent	85% ~ 90%

10 (O.D. = 2.0 at 1 micron film thickness and surface resistivity of 3.3×10^{11} ohm/square)

Method of Use (Application)

Photolithography process is applied to get fine resolution and wide β -bake window. Prime substrate is cleaned. 15 APX-K1, adhesion promoter from Shipley, is coated on substrate at 3000 rpm for 30 seconds, baked on hot plate at 175°C for 30 seconds. Black matrix formulation is coated on APX-K1 coated substrate at 750 rpm for 90 seconds, α -baked to evaporate solvent on hot plate at 100°C for 60 seconds. The 20 coatings were then β -baked in conventional ovens at 120°-180°C for 30 minutes. The polyamic acids is 30%-50% imidized in this process. Photoresist is coated at 5000 rpm for 30 seconds, soft baked on a hot plate at 100°C for 30 seconds, exposed and developed. Photoresist is stripped in safestrip. 25 Black matrix is then final cured in oven bake at 250°C for 30 minutes which completes the imidization process. Other colors are applied and processed.

Characteristic Properties

Figure 1 is a flow chart explaining the manufacturing 30 process of a black matrix according to the present invention.

Figure 2 is a graph showing transmission spectra of 1 μm film obtained in Example 1. The material meets the goals of optical density and high resistivity. Figure 3 shows sets of lithography data using both convection oven and hot plate beta-bake processes. Lithography results show a wide processing latitude. Resolution down to 3 μm at 1 μm film thickness are achievable within a wide range of intermediate (beta-bake) temperatures. Figure 4 is a schematic diagram illustrating photo of a resolution dagger. Figure 5 demonstrates Scanning Electron Microscope (SEM) of the black matrix material after resist removal and final cure. Good sidewall definition is evident in the pixel patterns. Figure 6 is a schematic diagram illustrating typical surface roughness measurement of the cured film. Surface roughness measurements were taken on a lithography test sample in an area adjacent to a resolution dagger. The surface has a uniform microroughness that is well suited to a black matrix application. Resistivity measurements for a 1 μm film are on the order of $10^{11} \Omega/\text{cm}^2$, thus providing a good balance of optical density and electrical properties. The composition can be applied by spin coating, imaged and developed using the same process used for RGB pixels. The average optical density for the 1 μm film thickness over the region 400-700 nm is 2.4. The material has good shelf life (3 months in freezer, 3 weeks at room temperature). It has low cost compared to chrome black matrix.

What is claimed:

1. A photolithographically imageable black matrix coating material having optical density ≥ 2.0 and surface resistivity greater than 10^5 ohm/square when coated at thicknesses ≤ 1.0 micron, consisting essentially of

5 a. a polyimide precursor vehicle and a solvent system therefore,

10 b. a soluble light-absorbing dye or mixtures of dyes, said dye or mixtures of dyes being substantially completely soluble in the vehicle and solvent system, and being effective to absorb substantially all light across a broad spectrum of from ultra violet to infrared, and

15 c. a mixed metal oxide non-carbon black pigment or mixture of pigments, and a dispersant therefor in Newtonian dispersion, said pigment or mixtures of pigments being substantially black in color.

2. The black matrix coating material of Claim 1, wherein the polyimide vehicle solvent system is an effective mixture of NMP and cyclohexanone, the dye mixture is Orasol brown 6RLN, and Orasol blue GN at a weight/weight ratio of 1:3 to 1:5 and the mixed metal oxide pigments are selected from the group consisting of spinel structured oxides of copper, manganese, chromium, iron, magnesium, aluminum, zinc, tin, titan, nickel, cobalt or mixtures thereof.

25 3. The black matrix coating material of Claim 1 having a dye specified from color index of solvent black 3 to solvent black 47.

4. The coating material of Claim 1 having a pigment specified from C.I. 77428 to 77494.

5. The coating material of Claim 1 wherein the polyimide vehicle is selected from the group of reaction products of

- a. ODA and PMDA, and
- b. ODA and BTDA.

6. The black matrix coating material of Claim 2 having a weight per cent composition of 4% to 8% polyimide vehicle, 10 50% to 80% solvent, 85% to 13% dye plus pigment wherein the weight/weight ratio of dye to pigment is 1:15 to 3:15, and the pigment is pigment black 26 made from a mixed metal oxide of copper, manganese and iron, having a C.I. of 77494, and 0.3% to 0.8% of a cationic dispersant made from high molecular 15 weight blocked copolymers with chemical groups having strong affinity to the pigment black 26.

7. The coating material of Claim 6 having a resistivity of at least 10¹¹ ohm/square.

8. The coating material of Claim 1 wherein the pigment 20 dispersion comprises pigment particles having a particle size less than 100 nm.

9. The coating material of Claim 1 wherein the surface of the pigments is coated with a material selected from the group consisting of silica, alumina and zirconia.

25 10. A method for making an organic black matrix coating material for STN or TFT pixel applications, said method comprising admixing

a. a polyimide precursor vehicle and light-absorbent organic broad spectrum light-absorbing dye capable of forming a high resistivity black coating and an effective solvent, and

5 b. a Newtonian dispersion of a black pigment having a C.I. from 77428 to 77494 at particle sizes less than 100 nm, and composed of mixed metal oxides of copper, manganese and iron, with an effective dispersant in the said effective solvent, and said dye and pigment being in a weight/weight
10 ratio of about 1:15 to 3:15,

whereby the coating materials optical density is higher than 2.0 at ultra thin coating thicknesses without the need for admixing carbon black.

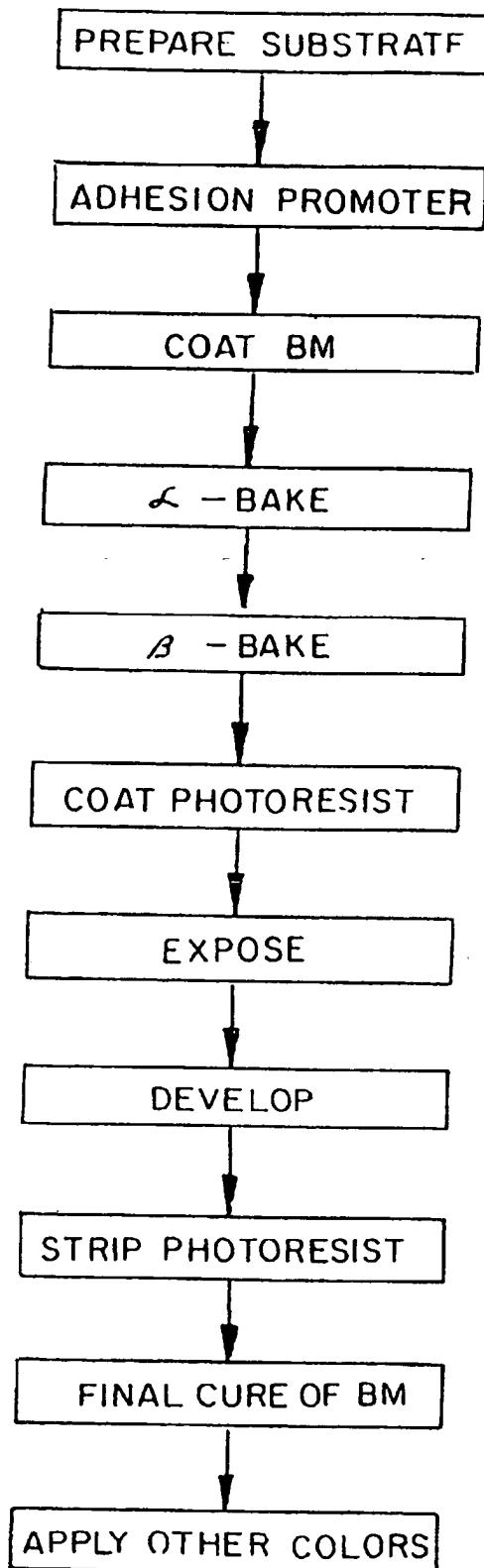


FIG. 1

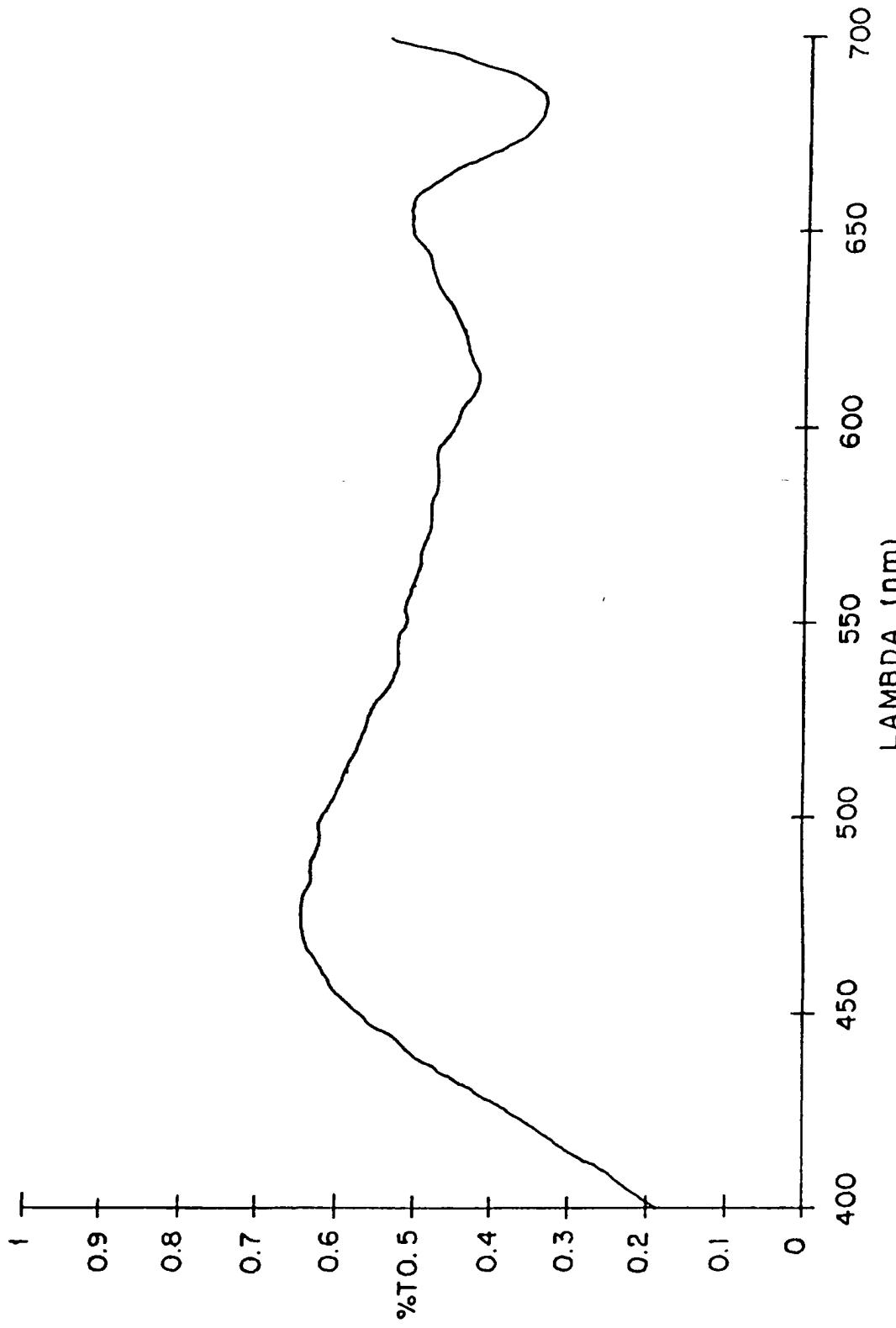


FIG.2

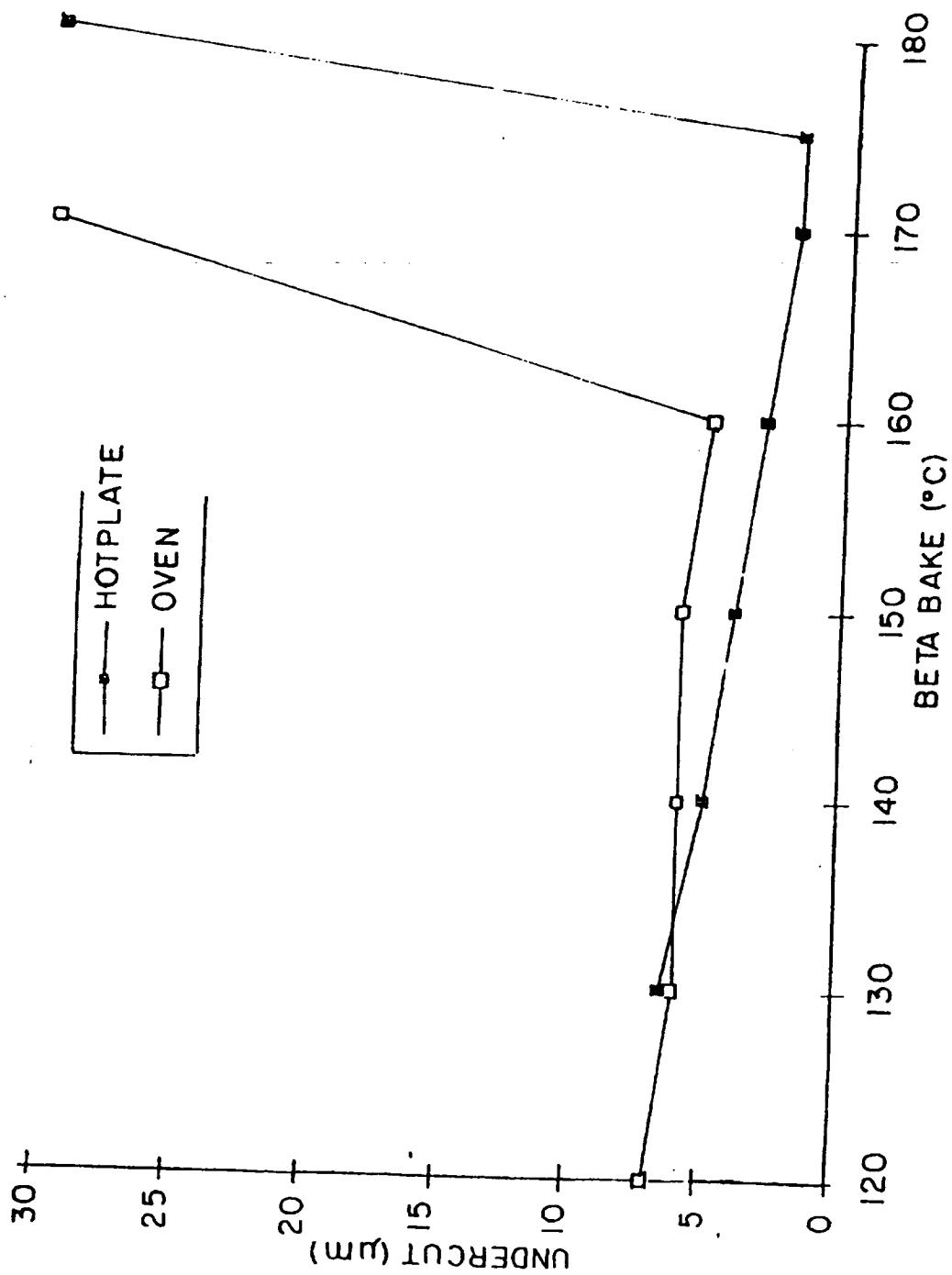


FIG. 3

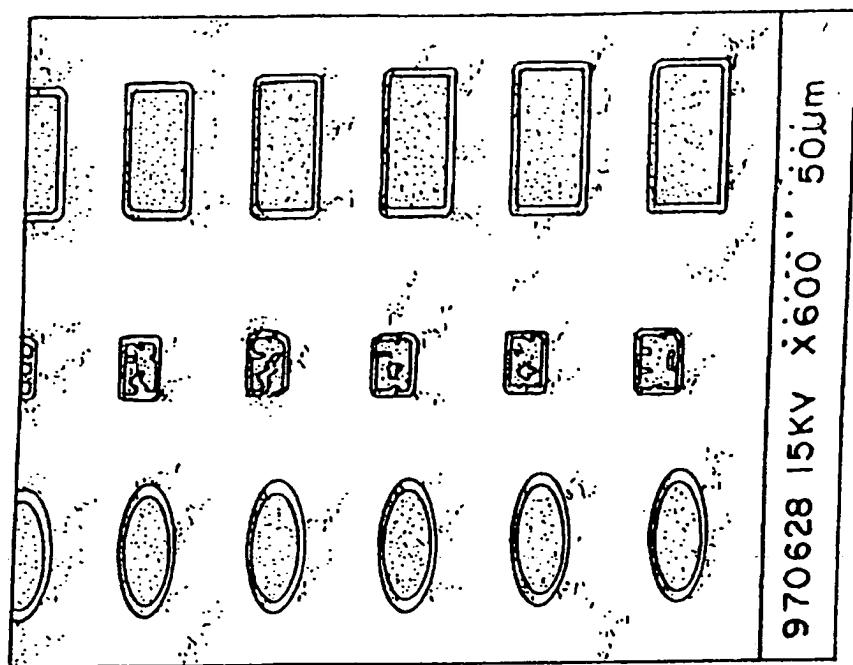


FIG.5

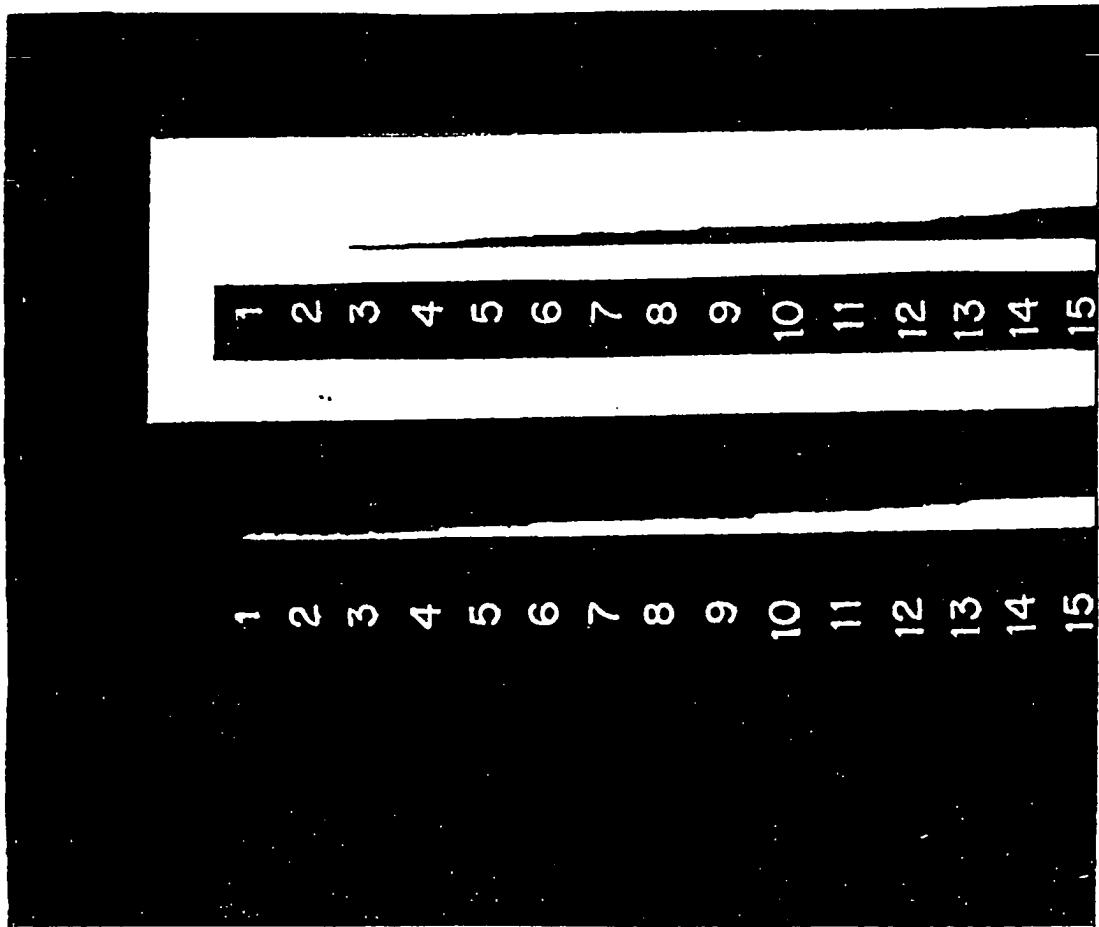


FIG.4

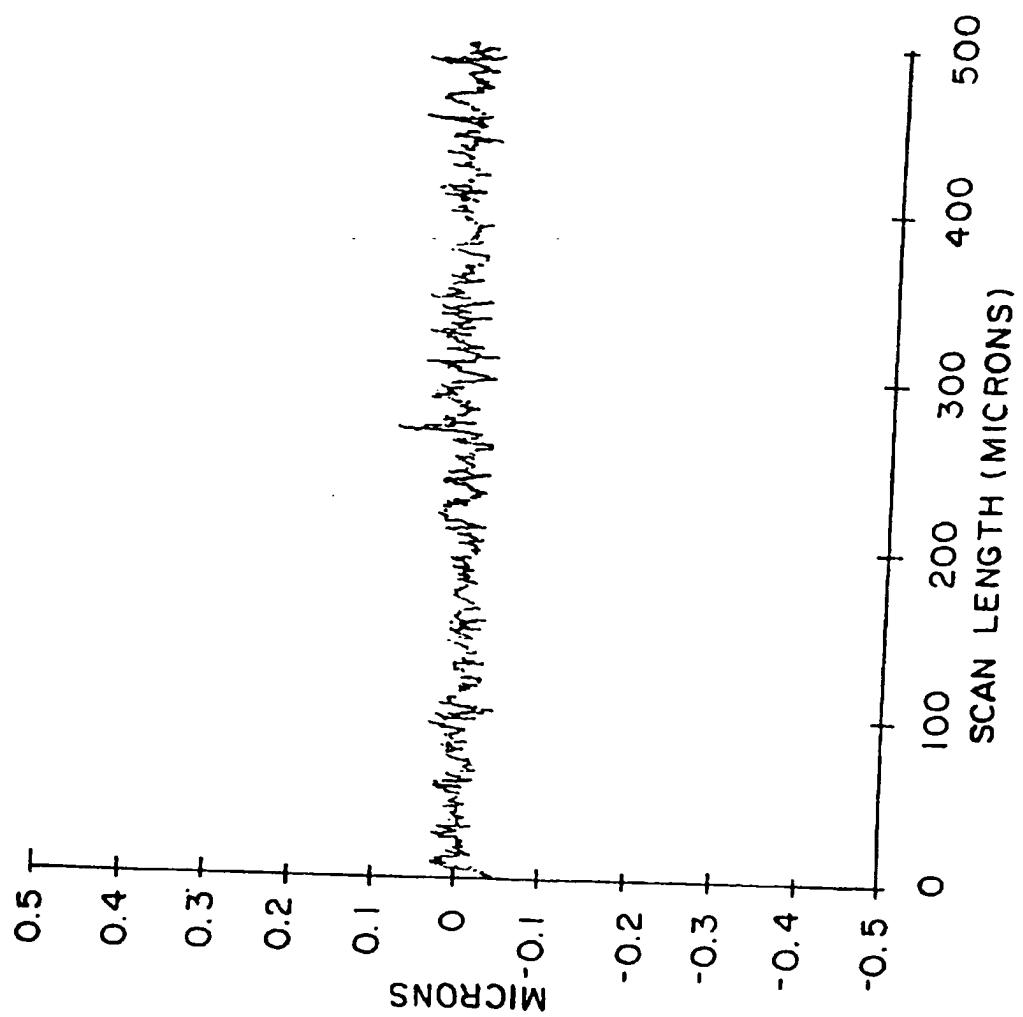


FIG. 6

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US97/11638

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) :G03C 1/73; G02B 5/20

US CL :430/270.1, 283.1, 7; 349/110

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 430/270.1, 283.1, 7; 349/110

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

APS

search terms: polyimide precursor, polyamic, black, dye, pigment, oxide

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 4,822,718 A (LATHAM ET AL) 18 April 1989, column 2, lines 13-27 and column 4, lines 7-42.	1-10
A	US 5,527,649 A (SATO ET AL) 18 June 1996, column 9, lines 5-42 and column 16, lines 12-41.	1-10
A, P	US 5,619,357 (ANGELOPOULOS ET AL) 08 April 1997, column 2, lines 18-55 and column 10, lines 13-31.	1-10

<input type="checkbox"/>	Further documents are listed in the continuation of Box C.	<input type="checkbox"/>	See patent family annex.
*	Special categories of cited documents:	"T"	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A"	document defining the general state of the art which is not considered to be of particular relevance	"X"	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"E"	earlier document published on or after the international filing date	"Y"	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"L"	document which may throw doubt on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Z"	document member of the same patent family
"O"	document referring to an oral disclosure, use, exhibition or other means		
"P"	document published prior to the international filing date but later than the priority date claimed		

Date of the actual completion of the international search	Date of mailing of the international search report
15 AUGUST 1997	23 OCT 1997
Name and mailing address of the ISA/US Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231 Facsimile No. (703) 305-3230	Authorized officer  JOHN MCPHERSON Telephone No. (703) 308-0661